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We claim:

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- 1. A fusion protein consisting essentially of an NS3, an NS4, and an NS5a polypeptide of a hepatitis C virus (HCV).
- A fusion protein consisting essentially of an NS3, an NS4, an NS5a, and NS5b polypeptide of an HCV.
- 3. A fusion protein according to either of claims 1 or 2, wherein one of the HCV polypeptides is derived from a different strain of HCV than the other HCV polypeptides.
 - 4. The fusion protein of claim 3 wherein each of the HCV polypeptides is derived from a different strain of HCV.
 - 5. A composition comprising:
 - (a) a fusion protein according to either of claims 1 or 2; and
 - (b) a pharmaceutically acceptable excipient.
- 20 6. A composition comprising:
 - (a) a fusion protein according to claim 4; and
 - (b) a pharmaceutically acceptable excipient.
 - 7. A composition consisting essentially of:
 - (a) an isolated and purified NS3 polypeptide of a hepatitis C virus (HCV);
 - (b) an isolated and purified NS4 polypeptide of a HCV;
 - (c) an isolated and purified NS5a polypeptide of a HCV; and
 - (d) a pharmaceutically acceptable excipient and optionally an adjuvant.

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The composition of claim 15 wherein the polynucleotide is in a plasmid.

	17.	A composition consisting essentially of:
		(a) an isolated and purified polynucleotide encoding an NS3 polypeptide
	of a hepatitis	C virus (HCV);
		(b) an isolated and purified polynucleotide encoding an NS4 polypeptide
5	of a HCV;	
		(c) an isolated and purified polynucleotide encoding an NS5a polypeptide
	of a HCV; an	nd ·
		(d) a pharmaceutically acceptable excipient and optionally an adjuvant.
10	18.	The composition of claim 17 wherein the polynucleotide is DNA.
	19.	The composition of claim 18 wherein the polynucleotide is in a plasmid.
	20.	A composition consisting essentially of:
15		(a) an isolated and purified polynucleotide encoding an NS3 polypeptide
	of a hepatitis C virus (HCV);	
		(b) an isolated and purified polynucleotide encoding an NS4 polypeptide
	of a HCV;	
		(c) an isolated and purified polynucleotide encoding an NS5a polypeptide
20	of a HCV;	
		(d) an isolated and purified polynucleotide encoding an NS5b polypeptide
	of a HCV; ar	ad .
		(e) a pharmaceutically acceptable excipient and optionally an adjuvant.
25	21.	The composition of claim 20 wherein the polynucleotide is DNA.
	22.	The composition of claim 21 wherein the polynucleotide is in a plasmid.

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23. A method of activating T cells which recognize an epitope of an HCV polypeptide, comprising the step of:

contacting T cells with a fusion protein of either of claims 1 or 2, whereby a population of activated T cells recognizes an epitope of the NS3, NS4, NS5a, or NS5b polypeptides.

- 24. The method of claim 23 wherein the T cells are obtained from a mammal selected from the group consisting of a mouse, a baboon, a chimpanzee, and a human.
- 10 25. The method of claim 24 wherein the mammal is infected with an HCV.
 - 26. The method of claim 24 wherein the mammal is not infected with an HCV.
- T cells. The method of claim 23 wherein the population of T cells comprises CD4⁺
 - 28. The method of claim 23 wherein the population of T cells comprises CD8⁺ T cells.
 - 29. The method of claim 28 wherein the CD8⁺ T cells express interferon-γ.
 - 30. The method of claim 28 wherein the CD8⁺ T cells specifically recognize an epitope of an NS5a polypeptide.
 - 31. The method of claim 30 wherein the epitope is selected from the group consisting of the epitopes shown in SEQ ID NO:1 and SEQ ID NO:2.

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- 32. The method of claim 23 wherein the T cells comprise CD8⁺ and CD4⁺ T cells.
- 33. The method of claim 23 wherein the step of contacting further comprisescontacting the T cells with an adjuvant.
 - 34. The method of claim 23 wherein the fusion protein is provided by a polynucleotide encoding the fusion protein.
- 10 35. The method of claim 34 wherein the polynucleotide is DNA.
 - 36. The method of claim 34 wherein the polynucleotide is RNA.
 - 37. The method of claim 23 wherein the T cells are in a mammal.
 - 38. The method of claim 37 wherein the mammal is selected from the group consisting of a mouse, a baboon, a chimpanzee, and a human.
 - 39. The method of claim 37 wherein the mammal is infected with an HCV.
 - 40. The method of claim 37 wherein the mammal is not infected with an HCV.
- 41. A method of activating T cells which recognize an epitope of an HCV polypeptide, comprising the step of:

contacting T cells with a composition according to claim 7, whereby a population of activated T cells recognizes an epitope of the NS3, NS4, NS5a, or NS5b polypeptides.

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42. A method of activating T cells which recognize an epitope of an HCV polypeptide, comprising the step of:

contacting T cells with a composition according to claim 8, whereby a population of activated T cells recognizes an epitope of the NS3, NS4, NS5a, or NS5b polypeptides.

43. A method of activating T cells which recognize an epitope of an HCV polypeptide, comprising the step of:

contacting T cells with a composition according to claim 17, whereby a population of activated T cells recognizes an epitope of the NS3, NS4, NS5a, or NS5b polypeptides.

44. A method of activating T cells which recognize an epitope of an HCV polypeptide, comprising the step of:

contacting T cells with a composition according to claim 20, whereby a

population of activated T cells recognizes an epitope of the NS3, NS4, NS5a, or NS5b polypeptides.